

What is claimed is:

1. A method of amplifying nucleic acid encoding at least a portion of an antibody comprising:

a) annealing a primer to a template that encodes at least a portion of an antibody,

5 the primer having a first portion which anneals to the template and a second portion of predetermined sequence which does not anneal to the template;

b) synthesizing a polynucleotide that is complementary to the portion of the template between the location at which the first portion of the primer anneals to the template and the end of the template, the polynucleotide having the primer at a first end

10 thereof and a second end;

c) separating the polynucleotide synthesized in step (b) from the template;

d) annealing a template oligonucleotide to the second end of the polynucleotide synthesized in step (b), the template oligonucleotide having a first portion that anneals to the second end of the polynucleotide and a second portion having the same

15 predetermined sequence as the second portion of the primer;

e) extending the polynucleotide synthesized in step (b) to provide a terminal portion thereof that is complementary to the predetermined sequence; and

f) amplifying the extended polynucleotide using a single primer having the predetermined sequence.

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2. A method as in claim 1 wherein the step of annealing a primer to a template that encodes at least a portion of an antibody comprises annealing at least one primer that comprises a sequence selected from the group consisting of

CTCGAGCAGGTCAGCTGGTGCAG (SEQ ID NO 296),
 CTCGAGCAGGTCCAGCTTGTGCAG (SEQ ID NO 297),
 CTCGAGSAGGTCCAGCTGGTACAG (SEQ ID NO 298),
 CTCGAGCARATGCAGCTGGTGCAG (SEQ ID NO 299),
 5 CTCGAGCAGATCACCTTGAAGGAG (SEQ ID NO 300),
 CTCGAGCAGGTCACCTTGARGGAG (SEQ ID NO 301),
 CTCGAGGARGTGCAGCTGGTGGAG (SEQ ID NO 302),
 CTCGAGCAGGTGCAGCTGGTGGAG (SEQ ID NO 303),
 CTCGAGGAGGTGCAGCTGTTGGAG (SEQ ID NO 304),
 10 CTCGAGCAGSTGCAGCTGCAGGAG (SEQ ID NO 305),
 CTCGAGCAGGTGCAGCTACAGCAG (SEQ ID NO 306),
 CTCGAGGARGTGCAGCTGGTGCAG (SEQ ID NO 307),
 CTCGAGCAGGTACAGCTGCAGCAG (SEQ ID NO 308) and
 CTCGAGCAGGTSCAGCTGGTGCAA (SEQ ID NO 309),
 15 wherein R is A or G, K is G or T, and S is C or G.

3. A method as in claim 1 wherein the step of annealing a primer to a template that encodes at least a portion of an antibody comprises annealing a primer to a template that encodes at least a portion of an IgA antibody.

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4. A method of amplifying nucleic acid encoding at least a portion of an antibody comprising:

a) annealing a primer and a boundary oligonucleotide to a template that encodes at least a portion of an antibody, the primer having a first portion which anneals to the

template and a second portion of predetermined sequence which does not anneal to the template;

b) synthesizing a polynucleotide that is complementary to the portion of the template between the location at which the first portion of the primer anneals to the template and the portion of the template to which the boundary oligonucleotide anneals, the polynucleotide having the primer at a first end thereof and a second end;

c) separating the polynucleotide synthesized in step (b) from the template;

d) annealing a template oligonucleotide to the second end of the polynucleotide synthesized in step (b), the template oligonucleotide having a first portion that anneals to the second end of the polynucleotide and a second portion having the same predetermined sequence as the second portion of the primer;

e) extending the polynucleotide synthesized in step (b) to provide a terminal portion thereof that is complementary to the predetermined sequence;

f) amplifying the extended polynucleotide using a single primer having the predetermined sequence.

5. A method as in claim 4 wherein the step of annealing a primer to a template that encodes at least a portion of an antibody comprises annealing at least one primer that comprises a sequence selected from the group consisting of

CTCGAGCAGGTCAGCTGGTGCAG (SEQ ID NO 296),

CTCGAGCAGGTCCAGCTTGTGCAG (SEQ ID NO 297),

CTCGAGSAGGTCCAGCTGGTACAG (SEQ ID NO 298),

CTCGAGCARATGCAGCTGGTGCAG (SEQ ID NO 299),

CTCGAGCAGATCACCTTGAAGGAG (SEQ ID NO 300),
CTCGAGCAGGTCACCTTGARGGAG (SEQ ID NO 301),
CTCGAGGARGTGCAGCTGGTGGAG (SEQ ID NO 302),
CTCGAGCAGGTGCAGCTGGTGGAG (SEQ ID NO 303),
5 CTCGAGGAGGTGCAGCTGTTGGAG (SEQ ID NO 304),
CTCGAGCAGSTGCAGCTGCAGGAG (SEQ ID NO 305),
CTCGAGCAGGTGCAGCTACAGCAG (SEQ ID NO 306),
CTCGAGGARGTGCAGCTGGTGCAG (SEQ ID NO 307),
CTCGAGCAGGTACAGCTGCAGCAG (SEQ ID NO 308) and
10 CTCGAGCAGGTSCAGCTGGTGCAA (SEQ ID NO 309),
wherein R is A or G, K is G or T, and S is C or G.

6. A method as in claim 4 wherein the step of annealing a primer to a template
that encodes at least a portion of an antibody comprises annealing a primer to a template
15 that encodes at least a portion of an IgA antibody.

7. A method of producing an antibody library comprising:

a) providing a diverse population of templates that encode at least a portion of an
IgA antibody;

20 b) contacting the diverse population of templates with at least one primer, the at
least one primer having a first portion which anneals to the templates and a second
portion of predetermined sequence which does not anneal to the templates;

c) synthesizing polynucleotides that are complementary to the portion of the
templates between the location at which the first portion of the primer anneals to the

template and the end of the templates, the polynucleotides having the primer at a first end thereof and a second end;

d) separating the polynucleotides synthesized in step (c) from the templates;

e) annealing at least one template oligonucleotide to the second end of the

5 polynucleotides synthesized in step (c), the at least one template oligonucleotide having a first portion that anneals to the second end of the polynucleotides and a second portion having the same predetermined sequence as the second portion of the primer;

f) extending the polynucleotides synthesized in step (c) to provide a terminal portion thereof that is complementary to the predetermined sequence; and

10 g) amplifying the extended polynucleotides using a single primer having the predetermined sequence.

8. A method as in claim 7 wherein the step of annealing a primer to a template that encodes at least a portion of an antibody comprises annealing at least one primer that

15 comprises a sequence selected from the group consisting of

CTCGAGCAGGTKCAGCTGGTGCAG (SEQ ID NO 296),

CTCGAGCAGGTCCAGCTTGTGCAG (SEQ ID NO 297),

CTCGAGSAGGTCCAGCTGGTACAG (SEQ ID NO 298),

CTCGAGCARATGCAGCTGGTGCAG (SEQ ID NO 299),

20 CTCGAGCAGATCACCTTGAAGGAG (SEQ ID NO 300),

CTCGAGCAGGTCACCTTGARGGAG (SEQ ID NO 301),

CTCGAGGARGTGCAGCTGGTGGAG (SEQ ID NO 302),

CTCGAGCAGGTGCAGCTGGTGGAG (SEQ ID NO 303),

CTCGAGGAGGTGCAGCTGTTGGAG (SEQ ID NO 304),

CTCGAGCAGSTGCAGCTGCAGGAG (SEQ ID NO 305),

CTCGAGCAGGTGCAGCTACAGCAG (SEQ ID NO 306),

CTCGAGGARGTGCAGCTGGTGCAG (SEQ ID NO 307),

5 CTCGAGCAGGTACAGCTGCAGCAG (SEQ ID NO 308) and

CTCGAGCAGGTSCAGCTGGTGCAA (SEQ ID NO 309),

wherein R is A or G, K is G or T, and S is C or G.

9. A library of IgA antibodies prepared in accordance with the method of claim

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10. A method of identifying an antibody having a desired binding specificity
comprising:

preparing a library of IgA antibodies in accordance with the method of claim 7;

and

15 screening the library to identify one or more IgA antibodies having a desired
binding specificity.

11. An IgA antibody identified in accordance with the method of claim 10.

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